

STIC-ILL

From: Borin, Michael
Sent: Thursday, February 11, 1999 3:50 PM
T : STIC-ILL
Subject: copy request

Examiner: M.Borin

AU: 1654

Tel.: 305-4506

Date needed: ASAP

_RE: 09/120030; lysostaphin

Please make copies of :

L22 ANSWER 55 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 29
AU OLDHAM E R; DALEY M J
SO J DAIRY SCI, (1991) 74 (12), 4175-4182.
CODEN: JDSCAE. ISSN: 0022-0302.

L22 ANSWER 71 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 35
AU BRAMLEY A J; FOSTER R
SO RES VET SCI, (1990) 49 (1), 120-121.
CODEN: RV TSA9. ISSN: 0034-5288.

L22 ANSWER 74 OF 280 CAPLUS COPYRIGHT 1999 ACS
IN Blackburn, Peter; Pollack, June
SO S. African, 38 pp.
CODEN: SFXXAB

L22 ANSWER 92 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS
AU SEARS P M; SMITH B S; POLAK J; GUSIK S N; BLACKBURN P
SO EIGHTY-THIRD ANNUAL MEETING OF THE AMERICAN DAIRY SCIENCE ASSOCIATION,
EDMONTON, ALBERTA, CANADA, JUNE 26-29, 1988. J DAIRY SCI. (1988) 71 (SUPPL
1), 244.
CODEN: JDSCAE. ISSN: 0022-0302. P. 244.

L22 ANSWER 188 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 93
AU EASMON C S F; LANYON H; COLE P J
SO BR J EXP PATHOL, (1978) 59 (4), 381-385.
CODEN: BJ EPA5. ISSN: 0007-1021.

L22 ANSWER 224 OF 280 CAPLUS COPYRIGHT 1999 ACS
AU Zyagmunt, Walter A.; Tavormina, Peter A.
SO Fortschr. Arzneimittelforsch. (1972), 16, 309-33
CODEN: FAZMAE

L22 ANSWER 228 OF 280 BIOSIS COPYRIGHT 1999 BIOSIS DUPLICATE 105

- P376 Lysostaphin efficacy for treatment of *Staphylococcus aureus* intramammary infection. P.M. Sears*, B.S. Smith, J. Polak, and S.N. Gusik, and P. Blackburn, Cornell University, Ithaca, NY and Public Health Research Institute/Applied Microbiology, Inc., New York, NY.

Clone-derived lysostaphin was evaluated as to its bacteriocidal effect on *S. aureus* intramammary infections. *S. aureus* (Newbould-305) was eliminated from glands of guinea pigs 48 hrs post-infection by 125 mcg of lysostaphin in 14/16, 25 mcg in 5/8, 5 mcg in 5/10, 1 mcg in 0/1, and 0 mcg in 0/3. Glands infected with *S. aureus* at 48 hrs post-challenge in untreated guinea pigs persisted; however 3/25 control glands of treated guinea pigs cleared in response to treatment of the adjacent gland.

Somatic cell/ml in the guinea pig shifted from 10's $\times 10^3$ pre-infected glands to cell counts greater than 1.0×10^6 following *S. aureus* inoculation. Treatment with lysostaphin caused a neutrophilic shift in the treated gland to levels exceeding 100×10^6 accompanied by an increase in the adjacent non-treated gland but dropped sharply to pre-treatment level. The greatest response in control glands was observed in animals receiving 125 mcg which corresponded to 2/25 clearance of *S. aureus* in control glands.

The leukocytic response to intramammary treatment in the cow is similar to the guinea pig model described above. Somatic cell levels increased ten fold in *S. aureus* infected glands at the milking following treatment. Cell levels returned to pre-treatment levels or lower in subsequent milkings. A rise in leukocytes alone could not account for clearance of the infection.

- P377 The effect of a hydraulic milking device on milking rate, milk yield and transfer of bacteria between quarters in dairy cows. L. M. Rode*, D. S. Croy, R. C. Phillippe and K.-J. Chang. Agriculture Canada, Lethbridge, Alberta, and Alberta Agriculture, Lethbridge, Alberta.

Sixteen cows in midlactation were used in a crossover design, with two periods of 21 days, to determine the effectiveness of a hydraulic milking device (Hydrast*[®], Deosan Ltd., Northampton, U.K.). Milk yield was measured every 15 seconds until milk flow ceased. Milk yield was 5.1 and 4.8 kg ($P < 0.01$) after 120 seconds and 7.4 and 7.2 kg ($P < 0.01$) after 180 seconds, for Hydrast-milked (H) and control (C) cows, respectively. Total milking time was 311 and 317 seconds for H and C respectively, and unaffected by treatment. Total milk yield was lower ($P < 0.01$) for H than C cows (10.4 vs. 10.9 kg per day). There was a time \times treatment interaction ($P < 0.01$) for 120 and 180 second milk yield and total milking time. Cows adapted to the Hydrast device by milking faster. Transfer of a noninfectious *Rhizobium* marker bacteria was reduced but not prevented by the Hydrast device.

- P378 Influence of use of LDBSA or iodophor teat dip on staphylococcal prevalence and new *Staphylococcus aureus* infection rate. R.J. Harmon*, B.E. Langlois, K. Akers, W.L. Crist and R.W. Eichen. University of Kentucky, Lexington.

All cows in a university dairy herd ($N = 113$) were paired by breed, age, stage of lactation, and quarter infection status and randomly assigned to a group receiving either 1% iodophor (I) or 1.940 linear dodecyl benzene sulfonic acid (LDBSA) teat dip. Duplicate quarter samples were taken bimonthly over the 12 months completed. There was a slight decline in coagulase-negative *Staphylococcus* spp. prevalence in both groups but little difference between groups. *S. aureus* prevalence increased from 8.3% to 12.3% (of quarters) in the LDBSA group and declined from 8.7% to 3.0% in the I group. Number of new *S. aureus* infections for LDBSA and I were 45 and 7 and new infection (NI) rates (NI/100 cow-days) were .215 and .033. Although more *S. aureus* infected cows in the LDBSA group left the herd, more *S. aureus* infections appeared to be spontaneously eliminated from the I group during lactation. The NI rate for *S. aureus* in the LDBSA group was similar to that observed in published studies, but I was more effective in this herd in limiting NI by *S. aureus*.